CHAPTER 2.3.13.

BOVINE SPONGIFORM ENCEPHALOPATHY

Article 2.3.13.1.

The recommendations in this Chapter are intended to manage the human and animal health risks associated with the presence of the bovine spongiform encephalopathy (BSE) agent in cattle (Bos taurus and B. indicus) only.

The following *commodities* can be safely traded:

- 1) without BSE related restrictions and regardless of the BSE status of the country:
 - a) milk and milk products
 - <u>semen and *in vivo* derived cattle embryos collected and handled in accordance with the recommendations of the International Embryo Transfer Society:</u>
 - c) hides and skins (excluding hides and skins from the head);
 - <u>d)</u> <u>gelatin and collagen prepared exclusively from hides and skins (excluding hides and skins from the head):</u>
- <u>subject to the prescribed conditions relating to the BSE status of the cattle population of the exporting country or zone:</u>
 - a) cattle;
 - b) fresh meat and meat products,
 - c) gelatin and collagen prepared from bones;
 - d) tallow and tallow derivatives, and dicalcium phosphate.

Standards for diagnostic tests are described in the Terrestrial Manual.

Article 2.3.13.2.

The BSE <u>risk</u> status of the cattle population of a country or zone can only be determined on the basis of the following criteria:

- the outcome of a risk assessment identifying all potential factors for BSE occurrence and their historic perspective, in particular:
 - a) the potential for introduction and recycling of the BSE agent through consumption by cattle of meat and bone meal or greaves of ruminant origin;
 - b) importation of *meat and bone meal* or *greaves* potentially contaminated with a transmissible spongiform encephalopathy (TSE) or feedstuffs containing either;
 - c) importation of animals or embryos/oocytes (other than cattle embryos described in Article 2.3.13.8.) potentially infected with a TSE;

- d) epidemiological situation concerning all animal TSE in the country or zone;
- e) extent of knowledge of the population structure of cattle, sheep and goats in the country or zone;
- f) the origin and use of ruminant carcasses (including fallen stock), by products and slaughterhouse waste, the parameters of the rendering processes and the methods of animal feed manufacture;
- <u>the outcome of a risk assessment (which is reviewed annually), based on Section 1.3 of this Terrestrial Code, identifying all potential factors for BSE occurrence and their historic perspective:</u>
 - a) Release assessment

This comprises an assessment of the likelihood that a transmissible spongiform encephalopathy (TSE) agent has been introduced via the importation of the following commodities potentially contaminated with a TSE agent:

- i<u>) meat-and-bone meal or greaves;</u>
- ii) live animals;
- iii) animal feed and feed ingredients;
- iv) products of animal origin for human consumption.
- b) Exposure assessment

<u>This comprises an assessment of the likelihood of exposure of the BSE agent to susceptible animal species, through a consideration of the following:</u>

- i) epidemiological situation concerning all animal TSE agents in the country or zone;
- <u>ii) recycling and amplification of the BSE agent through consumption by cattle of meatand-bone meal or greaves of ruminant origin, or other feed or feed ingredients contaminated with these;</u>
- the origin and use of ruminant carcasses (including fallen stock), by-products and slaughterhouse waste, the parameters of the rendering processes and the methods of animal feed manufacture;
- <u>iv)</u> <u>implementation and enforcement of feed bans, including measures to prevent cross-</u> <u>contamination of animal feed;</u>
- 2) on-going awareness programme for veterinarians, farmers, and workers involved in transportation, marketing and slaughter of cattle to encourage reporting of all cases of neurological disease in adult cattle and-fallen.stock;
- compulsory notification and investigation of all cattle showing clinical signs <u>consistent</u> <u>compatible</u> with BSE;
- 4) a BSE surveillance and monitoring system with emphasis on risks identified in point 1) above, taking into account the guidelines in Appendix 3.8.4.; records of the number and results of investigations should be maintained for at least 7 years;
- 5) examination in an approved laboratory of brain or other tissues collected within the framework

of the aforementioned surveillance and monitoring system.

Standards for diagnostic tests are described in the Terrestrial Manual.

Article 2.3.13.3.

BSE free country or zone

The cattle population of a country or zone may be considered free of BSE should the following conditions be met:

- 1) a risk assessment, as described in point 1) of Article 2.3.13.2., has been conducted and it has been demonstrated that appropriate measures have been taken for the relevant period of time to manage any risk identified;
- 2) <u>a level of surveillance and monitoring which complies with the requirements of Appendix 3.8.4</u> is in place, and either:
 - a) there has been no case of BSE; and either:
 - i) the criteria in points 2) to 5) of Article 2.3.13.2. have been complied with for at least 7 years; or
 - ii) the criteria in point 3) of Article 2.3.13.2. have been complied with for at least 7 years and it has been demonstrated that for at least 8 years no *meat-and-bone meal* or *greaves* have been fed to ruminants;

OR

- b) all *cases* of BSE have been clearly demonstrated to originate directly from the importation of live cattle, and the affected cattle as well as, if these are females, <u>all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, <u>are permanently identified, and their movements controlled</u>, and when slaughtered or at <u>death, are completely destroyed</u> their last progeny born within 2 years prior to, or after, <u>clinical onset of the disease, if alive in the country or zone, have been slaughtered and completely destroyed</u>; and either:</u>
 - i) the criteria in points 2) to 5) of Article 2.3.13.2. have been complied with for at least 7 years; or
 - ii) the criteria in point 3) of Article 2.3.13.2. have been complied with for at least 7 years and it has been demonstrated that for at least 8 years no *meat-and-bone meal* or *greaves* have been fed to ruminants;

OR

- c) the last indigenous *case* of BSE was reported more than 7 years ago,
 - i) the criteria in points 2) to 5) of Article 2.3.13.2. have been complied with for at least 7 years; and
 - the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been banned and the ban has been effectively enforced for at least 8 years; and
 - iii) the affected cattle as well as:
 - <u>if these are females, all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, are permanently the country or zone.</u>

<u>identified, and their movements controlled</u>, and when slaughtered or at death, are completely destroyed, and

- all cattle which, during their first year of life, were reared with the affected cattle during their first year of life, and which investigation showed consumed the same potentially contaminated feed during that period, if alive in the country or zone, are permanently identified and their movements controlled, and when slaughtered or at death, are completely destroyed, or
- where the results of the investigation are inconclusive, all cattle born in the same herd as, and within 12 months of the birth of, the affected cattle, if alive in the country or zone, are permanently identified and their movements controlled, and when slaughtered or at death, are completely destroyed.

and when slaughtered or at death, are completely destroyed.

Article 2.3.13.4.

BSE provisionally free country or zone

The cattle population of a country or zone may be considered as provisionally free of BSE should the following conditions be met:

- 1) a risk assessment, as described in point 1) of Article 2.3.13.2., has been conducted and it has been demonstrated that appropriate measures have been taken for the relevant period of time to manage any risk identified;
- 2) <u>a level of surveillance and monitoring which complies with the requirements of Appendix 3.8.4</u> <u>is in place, and</u> either:
 - a) there has been no *case* of BSE; and either:
 - the criteria in points 2) to 5) of Article 2.3.13.2. are complied with, but have not been complied with for 7 years; or
 - ii) it has been demonstrated that for at least 8 years no *meat-and-bone meal* or *greaves* have been fed to ruminants, but the criteria in point 3) of Article 2.3.13.2. have not been complied with for 7 years;

OR

- b) all *cases* of BSE have been clearly demonstrated to originate directly from the importation of live cattle, and the affected cattle as well as, if these are females, <u>all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, <u>are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed, their last progeny born within 2 years prior to, or after, clinical onset of the disease, if alive in the country or zone, have been slaughtered and completely destroyed; and either:</u></u>
 - i) the criteria in points 2) to 5) of Article 2.3.13.2. are complied with, but have not been complied with for 7 years; or
 - ii) it has been demonstrated that for at least 8 years no *meat-and-bone meal* or *greaves* have been fed to ruminants, but the criteria in point 3) of Article 2.3.13.2. have not been complied with for 7 years.

Country or zone with a minimal BSE risk

The cattle population of a country or zone may be considered as presenting a minimal BSE risk should the country or zone comply with the following requirements:

- a risk assessment, as described in point 1) of Article 2.3.13.2., has been conducted and it has been demonstrated that appropriate measures have been taken for the relevant period of time to manage any risk identified;
- 2) <u>a level of surveillance and monitoring which complies with the requirements of Appendix 3.8.4</u> is in place, and

EITHER:

- a) the last indigenous *case* of BSE was reported more than 7 years ago, the criteria in points 2) to 5) of Article 2.3.13.2. are complied with and the ban on feeding ruminants with *meat-and-bone meal* and *greaves* derived from ruminants is effectively enforced, but:
 - i) the criteria in points 2) to 5) of Article 2.3.13.2. have not been complied with for 7 years; or
 - ii) the ban on feeding ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has not been effectively enforced for 8 years;

OR

- b) the last indigenous *case* of BSE has been reported less than 7 years ago, and the BSE incidence rate, calculated on the basis of indigenous *cases*, has been less than one two cases per million during each of the last four consecutive 12-month periods within the cattle population over 24 months of age in the country or zone (Note: For countries with a population of less than one million adult cattle, the maximum allowed incidence should be expressed in cattle-years.), and:
 - i) the ban on feeding ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been effectively enforced for at least 8 years;
 - ii) the criteria in points 2) to 5) of Article 2.3.13.2. have been complied with for at least 7 years;
 - iii) the affected cattle as well as:
 - if these are females, all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, <u>are permanently identified, and their movements controlled, and</u> when slaughtered or at death, are completely destroyed, and
 - all cattle which, during their first year of life, were reared with the affected cattle during their first year of life, and, which investigation showed consumed the same potentially contaminated feed during that period, if alive in the country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed, or
 - where the results of <u>the</u> investigation are inconclusive, all cattle born in the same herd as, and within 12 months of the birth of, the affected cattle, <u>if alive in the</u>

<u>country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed.</u>

if alive in the country or zone, when slaughtered or at death, are completely destroyed.

Article 2.3.13.6.

Country or zone with a moderate BSE risk

The cattle population of a country or zone may be considered as presenting a moderate BSE risk if:

- 1) a risk assessment, as described in point 1) of Article 2.3.13.2., has been conducted, and the other criteria listed in Article 2.3.13.2. are complied with;
- 2) the BSE incidence rate has been <u>measured using a level of surveillance and monitoring which</u> <u>complies with the requirements of Appendix 3.8.4., and is:</u>
 - a) if based only on surveillance in accordance with Article 3.8.4.2., greater than or equal to, one indigenous *case* per million and less than or equal to, one hundred indigenous *cases* per million within the cattle population over 24 months of age in the country or zone calculated over the past 12 months; or
 - b) if based on surveillance in accordance with Articles 3.8.4.2., 3.8.4.3. and 3.8.4.4., greater than, or equal to, one two indigenous cases per million and less than, or equal to, two hundred indigenous cases per million within the cattle population over 24 months of age in the country or zone calculated over the past 12 months; or
 - c) less than one two indigenous cases per million for less than four consecutive 12-month periods (Note: For countries with a population of less than one million adult cattle, the maximum allowed incidence should be expressed in cattle-years.);
- 3) the affected cattle as well as:
 - a) if these are females, all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed, and
 - b) all cattle which, during their first year of life, were reared with the affected cattle during their first year of life, and, which investigation showed consumed the same potentially contaminated feed during that period, if alive in the country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed, or
 - c) where the results of <u>the</u> investigation are inconclusive, all cattle born in the same herd as, and within 12 months of the birth of, the affected cattle <u>if alive in the country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed.</u>

if alive in the country or zone, when slaughtered or at death, are completely destroyed.

Countries and zones where the BSE incidence rate has been less than one indigenous *case* per million within the cattle population over 24 months of age during each of the last four consecutive 12-month

periods, but where at least one of the other requirements to be considered as provisionally free from BSE or as presenting a minimal BSE risk is not complied with, shall be considered as countries or zones with a moderate BSE risk.

Article 2.3.13.7.

Country or zone with a high BSE risk

The cattle population of a country or zone may be considered as presenting a high BSE risk if it cannot demonstrate that it meets the requirements of another category.

Article 2.3.13.8.

Regardless of the BSE status of the *exporting country*, *Veterinary Administrations* should authorise without restriction the import or transit through their territory of the following *commodities*:

- 1) milk and milk products
- 2) semen and *in vivo* derived cattle embryos collected and handled in accordance with the recommendations of the International Embryo Transfer Society;
- 3) protein-free tallow (maximum level of insoluble impurities of 0.15% in weight) and derivatives made from this tallow:
- 4) dicalcium phosphate (with no trace of protein or fat);
- 5) hides and skins;
- 6) gelatin and collagen prepared exclusively from hides and skins.

Article 2.3.13.9.

When importing from a BSE free country or zone, *Veterinary Administrations* should require:

for all *commodities* from cattle not listed in Article 2.3.13.8.

the presentation of an *international veterinary certificate* attesting that the country or zone complies with the conditions in Article 2.3.13.3. to be considered as free of BSE.

Article 2.3.13.10.

When importing from a BSE provisionally free country or zone, *Veterinary Administrations* should require:

for cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.4. to be considered as provisionally free of BSE;
- cattle selected for export are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin and are not the progeny of BSE suspect or confirmed females.

Article 2.3.13.11.

When importing from a country or zone with a minimal BSE risk, *Veterinary Administrations* should require:

for cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.5. to be considered as presenting a minimal BSE risk;
- 2) the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been banned and the ban has been effectively enforced;
- 3) cattle selected for export:
 - a) are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin and are not the progeny of BSE suspect or confirmed females exposed cattle as described in point 2) b) iii) of Article 2.3.13.5.;
 - b) were born after the date from which the ban on the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been effectively enforced.

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Article 2.3.13.12.
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When importing from a country or zone with a moderate BSE risk, *Veterinary Administrations* should require:

for cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.6. to be considered as presenting a moderate BSE risk;
- 2) the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been banned and the ban has been effectively enforced;
- 3) cattle selected for export:
 - a) are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin and are not the progeny of BSE suspect or confirmed females exposed cattle as described in point 3) of Article 2.3.13.6;
 - b) were born after the date from which the ban on the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been effectively enforced.

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Article 2.3.13.13.
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When importing from a country or zone with a high BSE risk, *Veterinary Administrations* should require:

for cattle

the presentation of an *international veterinary certificate* attesting that:

1) the country or zone complies with the conditions in Article 2.3.13.7. to be considered as presenting a high BSE risk;

- 2) the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been banned and the ban has been effectively enforced;
- 3) all affected cattle as well as:
 - a) if these are females, all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, <u>are permanently identified, and their movements</u> <u>controlled, and</u> when slaughtered or at death, are completely destroyed, and
 - b) all cattle which, during their first year of life, were reared with the affected cattle during their first year of life, and, which investigation showed consumed the same potentially contaminated feed during that period, or
 - c) where the results of an the investigation are inconclusive, all cattle born in the same herd as, and within 12 months of the birth of, the affected cattle,

if alive in the country or zone, <u>are permanently identified</u>, <u>and their movements controlled</u>, <u>and</u> when slaughtered or at death, are completely destroyed;

- 4) cattle selected for export:
 - a) are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin and are not the progeny of BSE suspect or confirmed females;
 - b) were born at least 2 years after the date from which the ban on the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants was effectively enforced.

When importing from a BSE provisionally free country or zone, *Veterinary Administrations* should require:

for *fresh meat* (bone-in or deboned) and *meat products* from cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.4. to be considered as provisionally free of BSE;
- 2) ante-mortem inspection is carried out on all cattle from which the meat or *meat products* destined for export originate.

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Article 2.3.13.15.
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When importing from a country or zone with a minimal BSE risk, *Veterinary Administrations* should require:

for fresh meat (bone-in or deboned) and meat products from cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.5. to be considered as presenting a minimal BSE risk;
- 2) ante-mortem inspection is carried out on all cattle from which the meat or *meat products* destined for export originate;

- 3) cattle from which the meat or *meat products* destined for export originate were not subjected to a stunning process, prior to slaughter, with a device injecting compressed air or gas into the cranial cavity or to a pithing process (laceration, after stunning, of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity);
- 4) the *fresh meat* and *meat products* destined for export do not contain brain, eyes, spinal cord or mechanically separated meat from skull and vertebral column from cattle over 30 months of age, all of which have been removed in a hygienic manner completely removed in a manner to avoid contamination with these tissues.

Article 2.3.13.16.

When importing from a country or zone with a moderate BSE risk, *Veterinary Administrations* should require:

for fresh meat (bone-in or deboned) and meat products from cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.6. to be considered as presenting a moderate BSE risk;
- 2) the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been banned and the ban has been effectively enforced;
- 3) ante-mortem inspection is carried out on all bovines;
- 4) cattle from which the meat or *meat products* destined for export originate were not subjected to a stunning process, prior to slaughter, with a device injecting compressed air or gas into the cranial cavity or to a pithing process;
- 5) the *fresh meat* and *meat products* destined for export do not contain brain, eyes, spinal cord, distal ileum the tissues listed in point 1) of Article 2.3.13.19. nor mechanically separated meat from skull and vertebral column from cattle over 6 months of age, all of which have been removed in a hygienic manner completely removed in a manner to avoid contamination with these tissues.

Article 2.3.13.17.

When importing from a country or zone with a high BSE risk, *Veterinary Administrations* should require:

for fresh meat and meat products from cattle

the presentation of an *international veterinary certificate* attesting that:

- 1) the country or zone complies with the conditions in Article 2.3.13.7. to be considered as presenting a high BSE risk;
- 2) the meat destined for export does not contain the tissues listed in point 1) of Article 2.3.13.19., all of which have been removed in a hygienic manner completely removed in a manner to avoid contamination with these tissues;
- 3) the meat destined for export, if obtained from animals over 9 months of age, has been deboned and does not contain nervous and lymphatic tissues exposed during a deboning process, all of

- which have been removed in a hygienic manner completely removed in a manner to avoid contamination with these tissues;
- 4) the *meat products* destined for export are derived from deboned meat and do not contain the tissues listed in point 1) of Article 2.3.13.19. nor nervous and lymphatic tissues exposed during a deboning process, nor mechanically separated meat from skull and vertebral column of bovine animals, all of which have been removed in a hygienic manner completely removed in a manner to avoid contamination with these tissues;
- 5) a system is in operation enabling the *fresh meat* and *meat products* destined for export to be traced back to the *establishments* from which they are derived;
- 6) ante-mortem inspection is carried out on all bovines;
- 7) the cattle from which the *meat* or *meat products* destined for export originate:
 - a) were identified by a permanent identification system enabling them to be traced back to the dam and herd of origin;
 - b) are not the progeny of BSE suspect or confirmed females; and either:
 - i) were born after the date from which the ban on the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been effectively enforced; or
 - ii) were born, raised and had remained in herds in which no *case* of BSE had been confirmed for at least 7 years;
 - c) were not subjected to a stunning process, prior to slaughter, with a device injecting compressed air or gas into the cranial cavity or to a pithing process;
- 8) the feeding of ruminants with *meat-and-bone meal* and *greaves* derived from ruminants has been banned and the ban has been effectively enforced;
- 9) all affected cattle as well as:
 - a) if these are females, all their progeny born within 2 years prior to and after clinical onset of the disease, if alive in the country or zone, <u>are permanently identified, and their movements</u> <u>controlled, and</u> when slaughtered or at death, are completely destroyed, and
 - b) all cattle which, during their first year of life, were reared with the affected cattle during their first year of life, and, which investigation showed consumed the same potentially contaminated feed during that period, if alive in the country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed, or
 - c) where the results of an the investigation are inconclusive, all cattle born in the same herd as, and within 12 months of the birth of, the affected cattle, if alive in the country or zone, are permanently identified, and their movements controlled, and when slaughtered or at death, are completely destroyed.

if alive in the country or zone, when slaughtered or at death, are completely destroyed.

Article 2.3.13.18.

Ruminant-derived meat-and-bone meal or greaves, or any commodities containing such products, which

originate from countries with a minimal, moderate or high BSE risk should not be traded between countries.

Article 2.3.13.19.

- 1) From cattle of any age originating from a country or zone with a moderate or a high BSE risk, the following commodities, and any commodity contaminated by them, should not be traded for the preparation of food, feed, fertilisers, cosmetics, pharmaceuticals including biologicals, or medical devices: tonsils and intestine, and protein products derived from them. Food, feed, fertilisers, cosmetics, pharmaceuticals or medical devices prepared using these commodities should also not be traded.
- 2) From cattle originating from a country or zone with <u>a moderate or</u> a high BSE risk, that were at the time of slaughter over <u>6 12</u> months of age, the following commodities, and any commodity contaminated by them, should not be traded for the preparation of food, feed, fertilisers, cosmetics, pharmaceuticals including biologicals, or medical devices: brains, eyes, spinal cord, tonsils, thymus, spleen, intestines, dorsal root ganglia, trigeminal ganglia, skull and vertebral column and derived protein products derived from the preceding. Food, feed, fertilisers, cosmetics, pharmaceuticals or medical devices prepared using these commodities should also not be traded.

From cattle, originating from a country or zone with a moderate BSE risk, that were at the time of slaughter over 6 months of age, the following commodities, and any commodity contaminated by them, should not be traded for the preparation of food, feed, fertilisers, cosmetics, pharmaceuticals including biologicals, or medical devices: brains, eyes, spinal cord, distal ileum, skull, vertebral column and derived protein products. Food, feed, fertilisers, cosmetics, pharmaceuticals or medical devices prepared using these commodities should also not be traded.

3) From cattle, originating from a country or zone with a minimal BSE risk, that were at the time of slaughter over 30 months of age, the following commodities, and any commodity contaminated by them, should not be traded for the preparation of food, feed, fertilisers, cosmetics, pharmaceuticals including biologicals, or medical devices: brains, eyes and spinal cord, skull, vertebral column and derived protein products. Food, feed, fertilisers, cosmetics, pharmaceuticals or medical devices prepared using these commodities should also not be traded.

Article 2.3.13.20.

Veterinary Administrations of *importing countries* should require:

for gelatin and collagen prepared from bones and intended for food or feed, cosmetics, pharmaceuticals including biologicals, or medical devices

the presentation of an *international veterinary certificate* attesting that the bones came from:

- 1) a BSE free or provisionally free country or zone, or from a country or zone with a minimal BSE risk; or
- 2) a country or zone with a moderate BSE risk; and
 - a) skulls and vertebrae (excluding tail vertebrae) have been excluded;
 - b) the bones have been subjected to a process which includes all the following steps:

- i) pressure washing (degreasing),
- ii) acid demineralisation,
- iii) prolonged alkaline treatment,
- iv) filtration,
- v) sterilisation at ≥138°C for a minimum of 4 seconds,

or to an equivalent process in terms of infectivity reduction.

Veterinary Administrations of *importing countries* should require:

for tallow <u>and dicalcium phosphate</u> (other than protein free tallow as defined in Article 2.3.13.8.) intended for food, feed, fertilisers, cosmetics, pharmaceuticals including biologicals, or medical devices

the presentation of an *international veterinary certificate* attesting that it originates from:

- 1) a BSE free or provisionally free country or zone, or
- 2) a country or zone with a minimal BSE risk, and it originates from cattle which have been subjected to an ante-mortem inspection for BSE with favourable results and has not been prepared using the tissues listed in point 3 of Article 2.3.13.19., or
- 3) a country or zone with a moderate BSE risk, and it originates from cattle which have been subjected to an ante-mortem inspection for BSE with favourable results and has not been prepared using the tissues listed in point 2 of Article 2.3.13.19.

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Article 2.3.13.22.
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Veterinary Administrations of *importing countries* should require:

for tallow derivatives (other than those made from protein free tallow as defined in Article 2.3.13.8.) intended for food, feed, fertilisers, cosmetics, pharmaceuticals including biologicals, or medical devices

the presentation of an *international veterinary certificate* attesting that:

1) they originate from a BSE free or provisionally free country or zone, or from a country or zone with a minimal BSE risk;

OR

2) they have been produced by hydrolysis, saponification or transesterification using high temperature and pressure.

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Article 2.3.13.23.
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Careful selection of source materials is the best way to ensure maximum safety of ingredients or reagents of bovine origin used in the manufacture of medicinal products.

Countries wishing to import bovine materials for such purposes should therefore consider the

following factors:

- 1) the BSE status of the country and herd(s) where the animals have been kept, as determined under the provisions of Articles 2.3.13.2. to 2.3.13.7.;
- 2) the age of the donor animals;
- 3) the tissues required and whether or not they will be pooled samples or derived from a single animal.

Additional factors may be considered in assessing the risk from BSE, including:

- 1) precautions to avoid contamination during collection of tissues;
- 2) the process to which the material will be subjected during manufacture;
- 3) the amount of material to be administered;
- 4) the route of administration.

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APPENDIX 3.8.4.

SURVEILLANCE AND MONITORING SYSTEMS FOR BOVINE SPONGIFORM ENCEPHALOPATHY

Article 3.8.4.1.

Introduction

Surveillance for bovine spongiform encephalopathy (BSE) has at least two goals: to determine whether BSE is present in the country, and, if present, to monitor the extent and evolution of the epizootic, thus aiding control measures and monitoring their effectiveness.

The cattle population of a country or zone not free from BSE, will comprise the following sub-populations in order of decreasing size:

- 1) cattle not exposed to the infective agent;
- 2) cattle exposed but not infected;
- 3) infected cattle, which may lie within one of three stages in the progress of BSE:
 - a) the majority will die or be killed before reaching a stage at which BSE is detectable by current methods;
 - b) some will progress to a stage at which BSE is detectable by testing before clinical signs of disease appear;
 - c) the smallest number will show clinical signs of disease.

<u>A</u> surveillance programmes <u>on its own cannot guarantee BSE status and</u> should be determined by, and <u>be</u> commensurate with, the outcome of the risk assessment referred to in Article 2.3.13.2. and should take into account the diagnostic limitations associated with the above sub-populations and the relative distributions of infected animals among them.

Surveillance programmes developed before the advent of rapid diagnostic tests focused on the sub-population containing cattle displaying clinical signs compatible with BSE as described in Article 3.8.4.2. While Surveillance should focus on the sub-population containing cattle displaying clinical signs consistent with BSE as described in Article 3.8.4.2. this sub-population Where it is difficult to access all cattle displaying such clinical signs. investigation of other sub-populations using the new diagnostic techniques may provide a more accurate assessment picture of the BSE situation in the country or zone. A surveillance strategy programme may therefore need to combine several strategies. Recommended strategies for surveying the various sub-populations are described below.

Available data suggest the possibility that a gradient might be established to describe the relative value of surveillance applied to each sub-population. All countries should sample sub-populations identified in Articles 3.8.4.2. and 3.8.4.3. In countries where surveillance of cattle identified in Article 3.8.4.2. is unable to generate the numbers recommended in Table 1, surveillance should be enhanced by testing larger numbers of cattle identified in Article 3.8.4.3. Any shortfall in In addition, the first two sub-populations should be addressed by the surveillance can be complemented by sampling of normal cattle over 30 months of age at slaughter according to Article 3.8.4.4. Exclusive dependence on random sampling from normal cattle is not recommended, unless the number of

samples examined annually is statistically sufficient to detect a disease prevalence of 1 in 1,000,000.

Surveillance for BSE requires laboratory examination of samples in accordance with the methods described in the *Terrestrial Manual*.

For surveillance purposes, testing a part of the population is consistent with Chapter 1.3.6. on surveillance and monitoring of animal health.

Article 3.8.4.2.

Examination of cattle displaying clinical signs consistent with bovine spongiform encephalopathy

Cattle affected by illnesses that are refractory to treatment, and displaying progressive behavioural changes such as excitability, persistent kicking when milked, changes in herd hierarchical status, hesitation at doors, gates and barriers, as well as those displaying progressive neurological signs without signs of infectious illness are candidates for examination. Since BSE causes no pathognomonic clinical signs, all countries with cattle populations will observe individual animals displaying with compatible clinical signs consistent with BSE. It should be recognised that cases may display only some of these signs, which may also vary in severity, and such animals should still be investigated as potential BSE affected animals.

Table 1 indicates the minimum number of animals exhibiting one or more clinical signs of BSE that should be subjected to diagnostic tests according to the total cattle population over 30 months of age. The calculations assume a prevalence of one BSE clinically affected animal per one million adult cattle, a mortality rate not exceeding one percent per year in adult cattle, and a prevalence of central nervous system (CNS) signs not exceeding one percent within dying cattle.

As this sampling is not random, and as the mortality rate and prevalence of CNS signs within dying cattle may vary, the numbers indicated in this table are a subjective interpretation rather than a strict statistical deduction. This table should only be employed as a general guideline. Sampling in excess of the number indicated, ideally extending towards all cattle over 30 months of age showing clinical signs consistent with BSE, would give greater confidence in the outcome and is to be encouraged. In those cases where there is a shortfall in the number of samples required under this article, the difference may be made up by sampling in accordance with Article 3.8.4.3 and, in the event of a shortfall, by sampling in accordance with Article 3.8.4.4.

Table 1. Minimum number of annual investigations of cattle showing clinical signs consistent with BSE required for effective surveillance according to the total cattle population over 30 months of age

Total cattle population over 30 months of age	Minimum number of samples to examine	
500,000	50	
700,000	69	
1,000,000	99	
2,500,000	195	
5,000,000	300	
7,000,000	336	
10,000,000	367	

Total cattle population over 30 months of age	Minimum number of samples to examine
500,000	50
700,000	69
20,000,000	409
30,000,000	425
40,000,000	433

Article 3.8.4.3.

Examination of targeted cattle displaying clinical signs not necessarily indicative of bovine spongiform encephalopathy

Cattle <u>over 30 months of age</u> that have died or have been killed for reasons other than routine slaughter should be examined. This population will include cattle which have died on farm or in transit, 'fallen stock', and stock sent for emergency slaughter.

Many of these cattle may have exhibited some of the clinical signs listed in Article 3.8.4.2. which were not recognised as being compatible consistent with BSE. Experience in countries where BSE has been identified indicates that this population is the second most appropriate population to target in order to detect BSE. Empirical evidence indicates that surveillance conducted on one clinical suspect from Article 3.8.4.2. is equivalent to that conducted on 100 or more animals in this category in terms of its ability to detect BSE within an infected cattle population.

This multiplication factor of 100 should be applied in calculating the minimum sample size to substitute for any shortfall in the sample numbers specified in Article 3.8.4.2.

Article 3.8.4.4.

Examination of cattle subject to normal slaughter

In countries not free from BSE, sampling at routine slaughter of cattle over 30 months of age is a means of monitoring the progress of the epizootic and the efficacy of control measures applied, because it offers continuous access to a cattle population of known class, age structure and geographical origin. Empirical evidence indicates that surveillance conducted on one clinical suspect from Article 3.8.4.2. is equivalent to that conducted on 5,000 to 10,000 animals in this category in terms of its ability to detect BSE within an infected cattle population.

This multiplication factor of 5,000 to 10,000 should be applied in calculating the minimum sample size to substitute for any shortfall in the sample numbers specified in Article 3.8.4.2 and a multiplication factor of 50 to 100 applied regarding any shortfall in the sample numbers specified in Article 3.8.4.3.

Within each of the above sub-populations, countries may wish to target cattle identifiable as imported from countries or zones not free from BSE, cattle which have consumed potentially contaminated feedstuffs from countries or zones not free from BSE, offspring of BSE affected cows and cattle which have consumed feedstuffs potentially contaminated with other TSE agents.

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